

FORM PTO-1390 (REV. 1-2000)		U S DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE		ATTORNEY'S DOCKET NUMBER UMJ-105-B (UM1667)
TRANSMITTAL LETTER TO THE UNITED STATES DESIGNATED/ELECTED OFFICE (DO/EO/US) CONCERNING A FILING UNDER 35 U.S.C. 371				U.S. APPLICATION NO. (If known, see 37 CFR 1.5 09/936692
INTERNATIONAL APPLICATION NO. PCT/US00/06974	INTERNATIONAL FILING DATE 16 March 2000		PRIORITY DATE CLAIMED 17 March 1999	
TITLE OF INVENTION IONOMER COMPOSITE COMPOSITIONS				
APPLICANT(S) FOR DO/EO/US MA, Peter X. and HUBER, Anne E.				
Applicant herewith submits to the United States Designated/Elected Office (DO/EO/US) the following items and other information:				
<p>1. <input checked="" type="checkbox"/> This is a FIRST submission of items concerning a filing under 35 U.S.C. 371.</p> <p>2. <input type="checkbox"/> This is a SECOND or SUBSEQUENT submission of items concerning a filing under 35 U.S.C. 371.</p> <p>3. <input type="checkbox"/> This is an express request to begin national examination procedures (35 U.S.C. 371(f)). The submission must include items (5), (6), (9) and (21) indicated below.</p> <p>4. <input checked="" type="checkbox"/> The US has been elected by the expiration of 19 months from the priority date (Article 31).</p> <p>5. <input checked="" type="checkbox"/> A copy of the International Application as filed (35 U.S.C. 371(c)(2))</p> <ul style="list-style-type: none"> a. <input type="checkbox"/> is attached hereto (required only if not communicated by the International Bureau). b. <input type="checkbox"/> has been communicated by the International Bureau. c. <input checked="" type="checkbox"/> is not required, as the application was filed in the United States Receiving Office (RO/US). <p>6. <input type="checkbox"/> An English language translation of the International Application as filed (35 U.S.C. 371(c)(2)).</p> <ul style="list-style-type: none"> a. <input type="checkbox"/> is attached hereto. b. <input type="checkbox"/> has been previously submitted under 35 U.S.C. 154(d)(4). <p>7. <input checked="" type="checkbox"/> Amendments to the claims of the International Application under PCT Article 19 (35 U.S.C. 371(c)(3))</p> <ul style="list-style-type: none"> a. <input type="checkbox"/> are attached hereto (required only if not communicated by the International Bureau). b. <input type="checkbox"/> have been communicated by the International Bureau. c. <input type="checkbox"/> have not been made; however, the time limit for making such amendments has NOT expired. d. <input checked="" type="checkbox"/> have not been made and will not be made. <p>8. <input type="checkbox"/> An English language translation of the amendments to the claims under PCT Article 19 (35 U.S.C. 371(c)(3)).</p> <p>9. <input checked="" type="checkbox"/> An oath or declaration of the inventor(s) (35 U.S.C. 371(c)(4)).</p> <p>10. <input type="checkbox"/> An English language translation of the annexes of the International Preliminary Examination Report under PCT Article 36 (35 U.S.C. 371(c)(5)).</p>				
Items 11 to 20 below concern document(s) or information included:				
<p>11. <input checked="" type="checkbox"/> An Information Disclosure Statement under 37 CFR 1.97 and 1.98.</p> <p>12. <input checked="" type="checkbox"/> An assignment document for recording. A separate cover sheet in compliance with 37 CFR 3.28 and 3.31 is included.</p> <p>13. <input type="checkbox"/> A FIRST preliminary amendment.</p> <p>14. <input type="checkbox"/> A SECOND or SUBSEQUENT preliminary amendment.</p> <p>15. <input type="checkbox"/> A substitute specification.</p> <p>16. <input type="checkbox"/> A change of power of attorney and/or address letter.</p> <p>17. <input type="checkbox"/> A computer-readable form of the sequence listing in accordance with PCT Rule 13ter.2 and 35 U.S.C. 1.821 - 1.825.</p> <p>18. <input type="checkbox"/> A second copy of the published international application under 35 U.S.C. 154(d)(4).</p> <p>19. <input type="checkbox"/> A second copy of the English language translation of the international application under 35 U.S.C. 154(d)(4).</p> <p>20. <input type="checkbox"/> Other items or information:</p>				

17 SEP 2001

U.S. APPLICATION NO. (known as 37 CFR 1.492(e))
09/936692 INTERNATIONAL APPLICATION NO
 PCT/US00/06974

 ATTORNEY'S DOCKET NUMBER
 UMJ-105-B (UM 1667)

The following fees are submitted:

BASIC NATIONAL FEE (37 CFR 1.492 (a) (1) - (5)):

Neither international preliminary examination fee (37 CFR 1.482) nor international search fee (37 CFR 1.445(a)(2)) paid to USPTO and International Search Report not prepared by the EPO or JPO \$1000.00

International preliminary examination fee (37 CFR 1.482) not paid to USPTO but International Search Report prepared by the EPO or JPO \$860.00

International preliminary examination fee (37 CFR 1.482) not paid to USPTO but international search fee (37 CFR 1.445(a)(2)) paid to USPTO \$710.00

International preliminary examination fee (37 CFR 1.482) paid to USPTO but all claims did not satisfy provisions of PCT Article 33(1)-(4) \$690.00

International preliminary examination fee (37 CFR 1.482) paid to USPTO and all claims satisfied provisions of PCT Article 33(1)-(4) \$100.00

ENTER APPROPRIATE BASIC FEE AMOUNT =

CALCULATIONS PTO USE ONLY

Surcharge of \$130.00 for furnishing the oath or declaration later than 20 30 months from the earliest claimed priority date (37 CFR 1.492(e)).

\$ 100.00	
-----------	--

CLAIMS	NUMBER FILED	NUMBER EXTRA	RATE	\$
Total claims	31 - 20 =	11	x \$18.00	\$ 198.00
Independent claims	2 - 3 =		x \$80.00	\$
MULTIPLE DEPENDENT CLAIM(S) (if applicable)			+ \$270.00	\$

TOTAL OF ABOVE CALCULATIONS =

\$ 298.00	
-----------	--

Applicant claims small entity status. See 37 CFR 1.27. The fees indicated above are reduced by 1/2.

\$ 149.00	
-----------	--

SUBTOTAL =

\$ 149.00	
-----------	--

Processing fee of \$130.00 for furnishing the English translation later than 20 30 months from the earliest claimed priority date (37 CFR 1.492(f)).

\$	
----	--

TOTAL NATIONAL FEE =

\$ 149.00	
-----------	--

Fee for recording the enclosed assignment (37 CFR 1.21(h)). The assignment must be accompanied by an appropriate cover sheet (37 CFR 3.28, 3.31). \$40.00 per property +

\$ 40.00	
----------	--

TOTAL FEES ENCLOSED =

\$ 189.00	
-----------	--

Amount to be refunded:	\$
------------------------	----

charged:	\$
----------	----

- a. A check in the amount of \$ 189.00 to cover the above fees is enclosed.
- b. Please charge my Deposit Account No. _____ in the amount of \$ _____ to cover the above fees. A duplicate copy of this sheet is enclosed.
- c. The Commissioner is hereby authorized to charge any additional fees which may be required, or credit any overpayment to Deposit Account No. 25-0115. A duplicate copy of this sheet is enclosed.
- d. Fees are to be charged to a credit card. **WARNING:** Information on this form may become public. **Credit card information should not be included on this form.** Provide credit card information and authorization on PTO-2038.

NOTE: Where an appropriate time limit under 37 CFR 1.494 or 1.495 has not been met, a petition to revive (37 CFR 1.137 (a) or (b)) must be filed and granted to restore the application to pending status.

SEND ALL CORRESPONDENCE TO:

Julia Church Dierker
 YOUNG & BASILE, P.C.
 3001 West Big Beaver Rd.
 Suite 624
 Troy, Michigan 48084
 (248) 649-3333
 Fax: (248) 649-3338

SIGNATURE

 NAME
 Julia Church Dierker

REGISTRATION NUMBER
 33368

09/936692

Rec'd PCT/PTO 17 SEP 2001

PCT/US00/06974

WO 00/55253
6pt

IONOMER COMPOSITE COMPOSITIONS

CROSS REFERENCE TO RELATED APPLICATIONS

This application claims the benefit of U.S.

5 Provisional Application S.N. 60/124,819, filed March 17,
1999.

**STATEMENT REGARDING FEDERALLY SPONSORED
RESEARCH OR DEVELOPMENT**

10 This invention was made in the course of
research partially supported by a grant from the National
Institute of Health/National Institute of Dental Research
(NIH/NIDR) (Grant No. 2T32DE07057). The U.S. government
has certain rights in the invention.

BACKGROUND OF THE INVENTION

15 The present invention relates generally to
ionomer composite compositions, and more specifically to
such compositions useful for many dental applications,
ranging from direct restorative materials to fabrication
of preformed structure for dental and osseous tissue
repair applications.

20 It has been established that fluoride ions in
the oral environment are beneficial to the reduction of
recurrent caries, and formation of new caries. It is not
currently known why fluoride ions contribute to a
25 decrease in dental caries; theories include its effect on
the bacteria, the formation of fluorapatite, and an
increase in resistance to caries of both enamel and
dentin. The glass filler used in composites and glass
ionomers contains sufficient fluoride to see a reduction
30 in caries in filled and surrounding teeth. In addition,
the materials can be "recharged" through fluoride
toothpaste and topical fluoride treatments.

It has been shown that dental silicate cements
containing fluoride are therapeutic in preventing
35 secondary caries and reducing plaque formation. Acrylic
denture base material, restorative materials and
adhesives have been shown to be sites of bacterial and
plaque accumulation, which can be a precursor of

irritation to soft tissues and caries attack on remaining natural dentition. The release of fluoride ion from these dental restorative materials generally occurs either by surface release, or by dissolution of the
5 fluorine-containing additives or the dental restorative material itself with consequent migration of fluoride ions into the underlying tooth structure.

Various fluorine-containing additives that have been tried in dental restorations consist of inorganic fluoride salts, organic bases such as amine hydrofluoride, fluorocarbons and fluoride-containing ion-exchange resins. These attempts to find suitable fluorine-containing additives which are both dispersed in dental restorative material and capable of reducing tooth caries through controlled long-term fluoride release have failed. Silicate cements have demonstrated cariostatic release of fluoride. However, the strictly rapid surface release of fluoride from the cement, the dissolution of the cement in oral fluids, and the low tensile strengths of the cements are major disadvantages. Alternatively, the fluoride incorporated into insoluble resin materials has been considered to be virtually incapable of leaking out, and thus to be ineffective as a cariostatic agent. Studies by Forsten and Paunio (Scandinavian Journal of Dental Research (1972) 80: 515-519) comparing fluoride release by silicate cements and composite resins have shown that the overall release of fluoride from the two materials was comparable; however, the manner in which the fluoride was released from the composite was not controlled. It was found difficult to obtain controlled, effective cariostatic and plaque-reducing fluoride release from virtually insoluble materials such as acrylic denture base materials, adhesives and composite resins, and the like.

35 U.S. Patent No. 4,772,325 issued to Kwan et al. discloses a controlled, slow fluoride releasing additive comprising a Lewis base and a fluoride-containing Lewis

acid which is therapeutic in preventing secondary caries and reducing plaque formation. This additive is incorporated into polymeric dental restorative material and is capable of migrating from the interior to the surface of the material without dissolution thereof and with consequent release of fluoride. However, this composition is merely an additive to existing dental restorative materials. Further, it does not increase desirable mechanical properties of the dental restorative material in which it is incorporated; nor does it improve adhesion between polymers.

Current dental composite resins comprise one end of a continuous spectrum of multi-phase dental materials, with glass ionomers at the other end. Hybrid materials such as resin-modified glass ionomers and compomers lie somewhere between. Whereas both composite resins and glass ionomers have been known and used for a relatively long period of time, the hybrid materials such as the resin-modified glass ionomers and compomers are relatively new.

Ceramic filler reinforced polymeric composites are widely used in the area of dental restorative materials. A typical dental composite is composed of a mixture of silicate glass or quartz particles with an acrylic monomer that is polymerized to form a hardened composite material. In current dental composites, the fillers are mostly glasses, occasionally glass-ceramics and quartz (a crystalline form of silica), particulate polymers and glass-polymer composite particulates.

Strategies such as the development of smaller filler particles such as microfiller and nanofiller particles; the improvement of glass compositions; and the increase of filler volume fraction through the use of hybrid and heterogeneous filler systems have each been attempted in order to improve dental composites.

There is an increasing need to extend the use of polymeric composites to stress bearing posterior

applications due to concerns about the release of mercury from dental amalgam. However, the relatively low toughness, strength, wear resistance and durability of current dental composites have limited their use. It is generally agreed that current polymeric composites cannot be routinely substituted for dental amalgam and achieve the same clinical results. In the posterior dentition, in situations where occlusal stresses are concentrated, the current composites are inappropriate choices (Corbin and Kohn, 1994, JADA 125: 381-388). The current polymeric composites are not recommended for large posterior restorations because of the potential for excessive wear, microleakage or fracture (Bayne et al., 1994, JADA 125: 687-701). Composite restorations, in low stress-bearing applications not involving cusps, have average lifetimes of less than 10 years. In comparison, dental amalgam restorations, in high stress-bearing posterior applications with cusp replacement, have lifetimes of 15 years (Corbin and Kohn, 1994).

Many changes have occurred in recent years concerning dental restorative materials. Among these changes is the use of fluoride-releasing glass ionomer materials. Glass ionomer materials are based on the acid-base reaction of an aqueous solution of a polycarboxylic acid with an ion leachable, fluoride-containing glass (Wilson and Kent, 1971, J. Appl. Chem. Biotechnol. 21: 313; Wilson and Kent, 1972, Br. Dent. J. 132: 133). Glass ionomers are noted for their inherent adhesiveness to teeth and their ability to release fluoride to adjacent tooth structure in a sustained fashion to combat secondary caries.

However, it is generally accepted that glass ionomers possess inferior mechanical properties, including extreme brittleness and low strength (e.g., flexural strength of 10-20 MPa--McLean, 1990, J. Am. Dent. Assoc. 120: 43). This mechanical inferiority has severely limited their use. The reinforcement of glass

ionomers by disperse phase corundum (Prosser et al., 1986, J. Dent. Res. 65: 146), alumina fibers and other fibers (Scd and Wilson, Br. Pat. Application GB 2,028,855A, 1978), and metal powders (McLean and Gasser, U.S. Pat. No. 4,527,979, 1985) resulted in only incremental improvement in mechanical properties. Flexural strength values of reinforced glass ionomers have rarely exceeded 50 MPa (Wilson and McLean, Glass-Ionomer Cement, 1988; McLean, J. Am. Dent. Assoc., 1990, 120: 43).

Resin-modified glass ionomers (Mitra, EP Application 323,120, 1988; U.S. Pat. No. 5,154,762, 1992), where compatible resins (e.g., 2-hydroxyethyl methacrylate, or HEMA) are used with the polyacids, are only slightly stronger than glass ionomers (e.g., flexural strength of 60 MPa--Poolthong et al., 1994, Dent. Mater. J. 13: 220; Hickel, 1996, Acad. Dent. Mater. Trans. 9: 105). It has been generally accepted that the most intractable problem is likely to be lack of strength and toughness (Wilson, A.D. and J.W. McLean, Glass Ionomer Cement, Quintessence Publ. Co. (1988)). For a further discussion of glass ionomers as used in the dental market for direct restorative materials, eg. as cure-in-mouth cements, see, for example, Wilson and McLean, (1988); Katsuyama, S., T. Ishikawa, et al., Eds., Glass Ionomer Dental Cement - The Materials and Their Clinical Use, Ishiyaka EuroAmerica Inc. (1993); and O'Brien, W., Dental Materials and Their Selection, Chicago, Quintessence Publ. Co. (1997).

Compomers have been developed recently. See, for example, U.S. Pat. No. 4,816,495 issued to Blackwell et al.; U.S. Pat. No. 5,367,002 issued to Huang et al.; and Blackwell et al., Acad. Dent. Mater. Trans., 1996, 9: 77. Compomers are basically hybrid, glass ionomer-composites modified in their resin phase by a carboxylic acid monomer and in their filler phase by the inclusion of an acid-reactive, ion-leachable glass. The name

compomer is derived by combining the two words composite and ionomer, and is intended to suggest a combination of composite and glass-ionomer technology. The liquid part of a compomer is a mixture of a dental resin monomer (such as UDMA, a urethane dimethacrylate) and a carboxylic acid monomer (e.g., TCB, the reaction product of butane tetracarboxylic acid with HEMA), with the resin being the major phase and TCB the minor phase. The filler part of a compomer is a mixture of dental silicate glass and reactive fluorosilicate glass particles, with the reactive glass being the minor phase.

In contrast to glass ionomers, compomers do not generally contain significant amounts of water. The sole initial curing reaction is radical induced polymerization of the acrylic resin monomer matrix. An acid-base reaction takes place between TCB and the ion leachable fluorosilicate glass only after water infuses the cured composite via exposure to oral fluids, which also causes the filling to release fluoride ions. Flexural strength values of 90-125 MPa have been reported for compomers (Hickel, 1996, Acad. Dent. Mater. Trans. 9: 105).

However, these strength values are still inferior to those of current dental amalgam (110-150 MPa) and composite resins (100-145 MPa) (Hickel, 1996).

Therefore, compomers are currently not recommended for use in large, stress-bearing posterior applications.

It has been generally recognized that glasses and glass-ceramics are among the weakest and most brittle materials to use as reinforcement fillers. Glass filler particles are sensitive to surface flaws produced during mixing, handling and wear. A crack in the resin matrix can easily cut through the reinforcing glass particles.

U.S. Patent No. 5,861,445 issued to Xu et al. further defined geometrical shapes of the filler particles as cause for failure. The previously known glass fillers were either spherical or of irregular shapes, with length-to-diameter ratio only slightly

larger than one. Xu et al. disclosed that this had at least two major short-comings. First, rounded filler particles at occlusal surfaces are susceptible to facile dislodgement from the resin matrix during wear with foods bolus, resulting in high wear rates. Second, if a crack is initiated in the composite, it can easily propagate around the filler particles, hence causing the reinforcing effect of the filler particles to be lost.

As an answer to these shortcomings, Xu et al. proposed the use of ceramic filler particles and whiskers and/or chopped fibers to reinforce polymeric dental composites so that there are substantially improved mechanical properties and enhanced clinical longevity compared to conventional (or currently used) materials. The elongated whiskers and chopped fibers were said to have high length-to-diameter ratio values to effectively bridge and resist micro-cracks. In addition, it was asserted that the whiskers were less likely to be dislodged out of the matrix during wear.

In the area of conventional preformed structure for dental and osseous tissue repair applications, there are several shortcomings inherent therein. For example, current plastic teeth offer low abrasion resistance, low crazing resistance, and low heat-distortion temperatures when compared to porcelain teeth. See, for example, Craig, R., Ed., Restorative Dental Materials, St. Louis, Mosby (1997). If these problems could be overcome, plastic denture teeth are already superior to porcelain teeth in toughness, natural feel, and ease of grinding and polishing (Craig 1997).

Thus, it is an object of the present invention to provide an ionomeric composite composition for dental application(s) which advantageously improves the wear properties of the composition in the dental application. It is a further object of the present invention to provide such a composition having a highly crosslinked structure, thereby advantageously improving the strength

and crazing resistance of the composition in the dental application. Still further, it is an object of the present invention to provide such a composition which utilizes the unique properties of glass ionomers by advantageously adapting them for use in preformed denture teeth and other preformed structure. It is yet another object of the present invention to provide an ionomer composite advantageously utilizing a copolymer as one component thereof. Yet still further, it is an object of the present invention to provide such a composition which may advantageously be varied to suit a wide variety of dental applications, ranging from direct restorative materials, to intermediary dental materials such as liners, bases, and luting cement, to preformed structure for dental and osseous tissue repair applications. Yet further, it is an object of the present invention to provide such a composition which may advantageously be used for implants and/or tissue scaffolding (growing natural tissue within/on a porous synthetic material). It is still further an object of the present invention to provide such a composition which advantageously provides continuous fluoride release.

SUMMARY OF THE INVENTION

The present invention addresses and solves the above-mentioned problems and meets the enumerated objects and advantages, as well as others not enumerated, by providing ionomer composite compositions useful for dental applications. The composition consists essentially of a glass material containing at least one of divalent cations and multivalent cations, and optionally containing fluoride; and at least one copolymer. The copolymer comprises at least one hydrophilic monomer containing acid functional groups adapted to react with the divalent cations and/or multivalent cations to form ionic crosslinks among polymer chains. The hydrophilic monomer is present in an amount sufficient to impart a desired degree of aqueous

solubility to the copolymer. The copolymer further comprises at least one hydrophobic monomer present in an amount sufficient to impart a desired degree of structural stability to the composite composition when exposed to an aqueous environment.

The present ionomer composite composition is particularly advantageous in that, by varying the ratio of hydrophilic monomer to hydrophobic monomer, the copolymer may vary from water soluble to water insoluble. Thus, the present ionomer composite composition may be used for a wide variety of dental applications, ranging from those requiring water soluble compositions, eg. direct restorative materials, to intermediary materials such as for example, liners, bases, and luting cement, to those requiring water insoluble compositions, eg. to fabricate preformed structure for dental and osseous tissue repair applications.

BRIEF DESCRIPTION OF THE DRAWINGS

Other objects, features and advantages of the present invention will become apparent by reference to the following detailed description and drawings, in which:

Fig. 1 is a graph showing compressive modulus data for an 80:20 PMMA-MAA copolymer having varying glass contents;

Fig. 2 is a graph showing Vickers hardness data for the glass filled 80:20 PMMA-MAA copolymers;

Fig. 3 is a graph showing yield strength data for the glass filled 80:20 PMMA-MAA copolymers;

Fig. 4 is a graph showing toughness data for the glass filled 80:20 PMMA-MAA copolymers;

Fig. 5 is a graph showing modulus data for PMMA-MAA copolymers having varying MAA percentages, with 0 weight% and 50 weight% glass;

Fig. 6 is a graph showing Vickers hardness data for PMMA-MAA copolymers having varying MAA percentages, with 0 weight% and 50 weight% glass;

Fig. 7 is a graph showing yield strength data for PMMA-MAA copolymers having varying MAA percentages, with 0 weight% and 50 weight% glass;

5 Fig. 8 is a graph showing toughness data for PMMA-MAA copolymers having varying MAA percentages, with 0 weight% and 50 weight% glass;

Fig. 9 is a graph showing fluoride release data for a 75:25 PMMA-MAA copolymer having 50 weight% glass; and

10 Fig. 10 is a graph showing fluoride release data for an 80:20 PMMA-MAA copolymer having from 0 weight% to 80 weight% glass.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

15 The present invention provides novel ionomer composite compositions fortuitously useful for a heretofore unexpectedly wide range of dental applications. Such applications may range from direct restorative materials, eg. cure-in-mouth cements, to intermediary materials such as for example, liners, bases and luting cement, to preformed structure for dental and osseous tissue repair applications.

20 It is also to be understood that the present inventive compositions may find uses outside of the dental field, such as in orthopedic applications. Further, the fluoride releasing agent may be eliminated, if desired, from the copolymer-glass ionomeric composition described further hereinbelow; and the ionomeric composition(s) may find use as structural materials.

30 The present novel and inventive composition does not lie on the established continuum (discussed hereinabove) between conventional glass ionomers, resin-modified glass ionomers, polyacid-modified composite resins (compomers), and composite resins. Rather, the 35 novel composite compositions of the present invention possess unique properties.

The composition consists essentially of a glass material containing at least one of fluoride, divalent cations and multivalent cations; and at least one copolymer. The copolymer comprises at least one hydrophilic monomer containing acid functional groups adapted to react with the divalent cations and/or multivalent cations to form ionic crosslinks among polymer chains. The hydrophilic monomer is present in an amount sufficient to impart a desired degree of aqueous solubility to the copolymer. The copolymer further comprises at least one hydrophobic monomer present in an amount sufficient to impart a desired degree of structural stability to the composite composition when exposed to an aqueous environment.

It is to be understood that any suitable glass particles/materials may be chosen which contain divalent or/and multivalent cations. The glass material(s) optionally may contain fluoride(s). In a preferred embodiment, the glass material is selected from the group consisting of SiO_2 , Al_2O_3 , AlF_3 , CaF_2 , NaF , Na_3AlF_6 , AlPO_4 , and mixtures thereof. If a combination of these glass materials is chosen, such materials may be advantageously combined at various ratios.

The glass to polymer weight ratio may be between about 1:99 and about 95:5. In a preferred embodiment, the glass to polymer weight ratio is between about 10:90 and about 90:10. Still more preferred, the glass to polymer weight ratio is between about 40:60 and about 85:15.

It is to be understood that the composition of the present invention may comprise a copolymer or copolymers of two or more types of monomers, provided that at least one monomer is hydrophilic and at least one monomer is hydrophobic.

In accordance with the present invention, it is to be understood that any hydrophilic monomer(s) may be chosen provided it contains the above-mentioned acid

T02/T50-2162926560

functional groups adapted to react with the glass material(s)' divalent cations and/or multivalent cations to form ionic crosslinks among polymer chains. The acid functional groups also may react with calcium ions in teeth, leading to a desirable increase in the composition's adhesion properties with respect to the teeth.

Some exemplary suitable acid-containing hydrophilic monomers include the following: 1) Monomers containing carboxylic acid: acrylic acid, methacrylic acid, 4-vinylbenzoic acid, crotonic acid, oleic acid, elaidic acid, itaconic acid, maleic acid, fumaric acid, acetylenedicarboxylic acid, tricarballylic acid, sorbic acid, linoleic acid, linolenic acid, eicosapentenoic acid, other unsaturated carboxylic acids, anhydrides, their derivatives, and mixtures thereof; 2) Other organic acids such as sulfonic acid, and/or phosphonic acid replacement of the carboxyl group of the above listed unsaturated carboxylic acids, their derivatives, and mixtures thereof.

In addition to the one or more hydrophobic monomers and the one or more acid-containing hydrophilic monomers in the present inventive co-polymer composition, other hydrophilic monomers that do not have a reactive carboxyl group may be used as a co-monomer within the present inventive composition. An example of such a non-acid-containing hydrophilic monomer is 2-hydroxyethyl methacrylate (HEMA).

It is to be understood that, in accordance with the present invention, any hydrophobic monomer may be chosen, provided it imparts the desired degree of structural stability to the composite composition when exposed to an aqueous environment.

Some exemplary suitable hydrophobic unsaturated monomers include the following. Acrylates, methacrylates (eg. methyl methacrylate), ethylene, propylene, tetra-fluoroethylene, styrene, vinyl chloride, vinylidene

chloride, vinyl acetate, acrylonitrile, 2,2-bis[4-(2-hydroxy-3-methacryloyloxy-propoxy)-phenyl] propane (Bis-GMA), ethyleneglycol dimethacrylate (EGDMA), triethyleneglycol dimethacrylate (TEGDMA), bis(2-methacryloyloxyethyl) ester of isophthalic acid (MEI), bis(2-methacryloyloxyethyl) ester of terephthalic acid (MET), bis(2-methacryloyloxyethyl) ester of phthalic acid (MEP), 2,2-bis(4-methacryloyloxy phenyl) propane (BisMA), 2,2-bis[4-(2-methacryloyloxyethoxy) phenyl] propane (BisEMA), 2,2-bis[4-(3-methacryloyloxy-propoxy) phenyl] propane (BisPMA), hexafluoro-1,5-pentanediol dimethacrylate (HFPDMA), bis-(2-methacryloyloxyethoxy-hexafluoro-2-propyl) benzene [Bis(MEHFP) ϕ], 1,6-bis(methacryloyloxy-2-ethoxycarbonylamino)-2,4,4-tri-methylhexan (UEDMA), spiro orthocarbonates, other vinyl monomers, the derivatives of these monomers, and mixtures thereof.

The ratio of the hydrophilic monomeric units to the hydrophobic monomeric units may vary between about 1:99 and about 99:1. In a preferred embodiment, the hydrophilic monomer to hydrophobic monomer ratio is between about 5:95 and 95:5. By varying the monomer ratio, the copolymer could vary from water soluble (suitable for direct restorative dental applications) to water insoluble (suitable for fabrication of preformed dental structure and the like). As the amount of hydrophilic monomer is increased, the copolymer (as well as the ionomer composite composition) becomes more water soluble.

It is to be understood that any combination of the above hydrophilic and hydrophobic monomers may be suitable to form the copolymer component according to the present invention. An exemplary list of some copolymers includes the following: poly(methyl methacrylate-methacrylic acid), poly(methyl acrylate-acrylic acid), poly(methyl methacrylate-acrylic acid), poly(ethyl acrylate-acrylic acid), poly(ethyl methacrylate-methacrylic acid), poly(butyl acrylate-acrylic acid),

poly(ethylene-acrylic acid), poly(ethylene-methacrylic acid), poly(acrylonitrile-maleic anhydride), poly(butadiene-acrylonitrile-acrylic acid), poly(butadiene-maleic acid), poly(butadiene-maleic anhydride), poly(acrylamide-acrylic acid), poly(2-hydroxyethyl methacrylate-methacrylic acid), poly(propylene-acrylic acid), poly(propylene-ethylene-acrylic acid), poly(vinyl chloride-vinyl acetate-maleic acid), and mixtures thereof.

It is to be understood that the copolymers in the ionomer composite systems of the present invention may be polymerized before, during or after the ionomer reactions (base-acid reactions). In a preferred embodiment, the polymers or copolymers are polymerized before the ionomer reaction in order to reduce the contents of unreacted monomers or oligomers to improve biocompatibility.

According to the present invention, there may be a mixture of one or more of the above-mentioned copolymers; and other polymers in minor amounts may be added to the ionomer composite systems of the present invention. Examples of some such polymers which may be present as minor components include, but are not limited to poly(methyl methacrylate) (PMMA), polycarbonates, polyethylenes, polyamides, poly(ether-ether ketone) (PEEK) polymers, epoxies, and mixtures thereof.

It is to be understood that the copolymers and/or polymers mentioned above may be obtained from a commercial source; or they may be synthesized from appropriate monomer(s). If they are synthesized, the monomer(s) are suitably polymerized. The synthesized polymer/copolymer is then preferably subjected to a purification step in order to remove any unreacted monomer(s). The copolymer and polymer(s) in minor amounts (if any) may then be mixed and reacted with glass particles as described further hereinbelow.

It is further contemplated as being within the purview of the present invention to include other minor components in the ionomer composite composition(s) of the present invention. For example, pigments, surfactants, adhesion enhancers, fluoride releasing enhancers, and bioactive agents, such as for example, growth factors and hormones, and mixtures thereof may be incorporated thereinto if desired.

According to the present invention, the glass particles and the copolymers are mixed and reacted to form ionomer composite materials/compositions. The fluoride-containing glass particles can be dissolved or partially dissolved in a mixture of water and organic solvent(s), or in an aqueous solution. The dissolved divalent and multivalent cations form ionic bonds with the acid groups on the copolymer to impart the system with the characteristics of a glass ionomer. The undissolved glass particles act as a reinforcing second phase. This fortuitous result stemming from the present novel ionomeric composite composition is in contrast to the generally accepted theory (discussed hereinabove) that glasses are too weak and brittle to provide satisfactory reinforcement to dental restorative materials.

When polymerizable monomers, oligomers, and initiators are involved, a reinforcing polymerization may be carried out after the glass ionomer reaction, either thermally or photo-chemically, depending on the initiator type. The mechanical properties such as strength, moduli such as shear modulus, compressive modulus and tensile modulus, and hardness may be adjustable by varying the polymer composition and glass content.

The ionomer composite materials of the present invention may advantageously be thermally molded or solution cast to fabricate preformed structure for dental and osseous tissue repair applications. In addition, the present inventive materials may also be used for

applications of traditional glass ionomers and resin composites. As such, the present inventive materials may be particularly adapted for use for inlays, onlays, artificial teeth, denture base, filling materials, cavity liner, adhesive, orthodontic resin, pit and fissure sealant, dental, orthopedic and bone implants.

Further advantages flow from the ion releasing properties, which may impart the inventive materials with desirable bioactivities. Of specific importance is the fluoride release that can prevent secondary caries and reduce plaque formation in dental applications.

A preferred embodiment useful for preformed dental structure comprises a novel glass ionomer: poly(methyl methacrylate-co-methacrylic acid) (PMMA-MAA) glass ionomer. This material differs from standard glass ionomers in that the copolymer, in certain hydrophilic monomer to hydrophobic monomer ratios, is not water soluble, thereby reducing undesirable reaction with the oral environment. It also differs from PMMA based ionomers (see, for example, Ma, X., J.A. Sauer, et al., "Poly(methyl methacrylate) Based Ionomers. 1. Dynamic Mechanical Properties and Morphology," Macromolecules 28(11): 3953-3962 (1995)) by utilizing glass as a reinforcing second phase and ions from the glass in the crosslinking.

It has been fortuitously found that, by employing a PMMA-MAA copolymer, both the advantages of the methyl methacrylate unit and the ionomer character of the methacrylic acid unit, work in combination to form a strong and soundly crosslinked structure.

The glass was chosen for its fluoride release capabilities. See, for example, Kent, B., B. Lewis, et al., "Glass Ionomer Cement Formulations: I. The Preparation of Novel Fluoroaluminocilicate Glasses High in Fluorine," Journal of Dental Research 58(6): 1607-1619 (1979); Wilson, A.D., S. Crisp, et al., "Aluminocilicate Glasses for Polyelectrolyte Cements," Ind. Eng. Chem.

Prod. Res. Dev. 19: 263-270 (1980); de Araujo, F.B., F. Garcia-Goody, et al., "Fluoride release from fluoride-containing materials," Operative Dentistry 21: 185-190 (1996); Forsten, L., "Fluoride release and uptake by glass-ionomers and related materials and its clinical effect," Biomaterials 19: 503-508 (1998); and Verbeeck, R.M.H., E.A.P. De Maeyer, et al., "Fluoride release process of (resin-modified) glass-ionomer cements versus (polyacid-modified) composite resins," Biomaterials 19: 509-519 (1998).

In a preferred embodiment, the polymeric component of the composite composition comprises a copolymer, as opposed to a monomer(s), oligomers, or a completely water soluble polymer(s). This is preferable to *in situ* polymerization of a monomer(s), in that unreacted monomers remaining in the patient's mouth may have a deleterious effect on the mechanical properties of the dental restorative material, as well as on the patient's health.

As shown below in the Examples, the addition of glass to the PMMA-MAA copolymer appears to increase the compressive modulus, yield strength, and hardness of the material. It is believed that the improvement is no longer significant above about 70 weight% glass due, at least in part, to a limit in the packing density.

It appears that samples containing a greater amount of methacrylic acid units possess better mechanical properties within the copolymer range studied below (up to 25%) due, at least in part, to the larger number of crosslinks.

Without being bound to any theory, it is believed that the ionic interactions, both the polymer-polymer crosslinking and the polymer-glass ionic bonding, contribute to the increase in mechanical properties of the PMMA-MAA glass ionomer material.

Further, as can be seen below, experimental data indicate that the PMMA-MAA glass ionomer material is capable of sustained fluoride release.

To further illustrate the composition, the following examples are given. It is to be understood that these examples are provided for illustrative purposes and are not to be construed as limiting the scope of the present invention.

These initial studies evaluate PMMA-MAA glass ionomers using both hardness and compression testing.

EXAMPLE 1

Glass Incorporation. Two sets of samples were prepared for testing. In the first group, an 80:20 PMMA-MAA random copolymer was used with the following glass contents: 0.00, 6.25, 14.29, 25.00, 33.33, 40.00, 45.45, 50.00, 60.53, 70.00, and 80.00 weight percent. The second set includes the following PMMA-MAA random copolymers: 75:25, 80:20, 95:5, and 100:0, containing 0 wt% and 50 wt% glass. The copolymers were obtained from Polysciences, Inc., Warrington, Pennsylvania, the PMMA from Sigma Chemical Co., St. Louis, Missouri, and the glass from Specialty Glass Inc., Oldsmar, Florida, with a composition of 25.10 wt% SiO₂, 37.45 wt% Al₂O₃, and 37.45 wt% CaF₂. The glass particles are irregular in shape, range from 1-20 microns, and have no surface treatment.

To incorporate the glass, the polymer was dissolved at a 1/12 g/ml ratio in a 20% methanol, 80% benzene solution. For the 75:25 copolymer it was necessary to use a 30:70 solvent mix. The polymer and solvent were placed in a vial and vibrated on a shaker overnight. After 24 hours, the glass and a stir bar were added, and the mixture was stirred for one hour to incorporate the glass and allow time for the glass ionomer reaction to occur. The sample was frozen in a -20°C freezer, then placed on a lyophilizer in ice/salt bath to remove the solvent. A liquid nitrogen solvent

trap was maintained for two days, and the samples allowed to stay on the lyophilizer for a total of seven days.

EXAMPLE 2

Compression Molding Process. The following protocol was used to make samples for hardness and compression testing. A 4 mm diameter compression mold was filled with from 0.2 g to 0.3 g of sample (obtained by the procedure of Example 1). The sample amount was varied due to the density difference between the filled and unfilled polymer. The mold was placed in an oven at 450°F for two hours, then moved to a preheated 420°F press. The plunger was pressed down with 2000 lbs. force which was applied for 45 minutes, being monitored every 15 minutes, and increased if it had fallen. The pressure was then released and the plunger removed. The sample was allowed to stay on the press at 420°F for another 45 minutes to remove any residual compressive stresses. At this time the heater was turned off, and the sample cooled to room temperature before being removed from the mold.

EXAMPLE 3

Compression Testing. The cylindrical samples prepared in the compression mold of Example 2 were ground so that the ends were parallel. All samples possessed a length to diameter (L/D) ratio of 2.5. The samples were compressed to failure using a compression jig on an Instron 8521 mechanical tester with a crosshead speed of 0.5 mm/min. Five specimens were tested for each data point.

EXAMPLE 4

Hardness Testing. Samples prepared as in Example 2 were placed in liquid nitrogen and fractured with a hammer. A fragment no less than 2 mm thick was then mounted in Kould Mount. These samples were ground to 600 grit silicon carbide followed by polishing with a one micron diamond paste. A Tukon Microhardness Tester, model LR, was used to produce and measure Vickers indents

using a 300g load. Three indents were made on each sample and averaged.

EXAMPLE 5

Ion Bonding Test. In order to ascertain the effect of ions leached from the glass in crosslinking the polymer, an additional test was done. Glass powder was placed in a 20:80 methanol/benzene solvent mix (a 30:70 solvent mix for the 75/25 copolymer) and stirred for one hour. The glass was removed by centrifuging for one minute and decanting off the "ion solvent". This "ion solvent" was then used to prepare 0% glass PMMA-MAA compression and hardness samples as described above for each of the copolymers.

EXAMPLE 6

Fluoride Release Test. Five glass ionomer samples 1 cm x 1 cm x 0.05 cm were made for all fourteen variations (see Example 7 below) of the material, including the different glass contents and the different copolymers containing 50 wt% glass. Each sample was placed in a vial containing 5 ml ddH₂O. These samples were maintained on a shaker at 37°C for the duration of the experiment. Each day the water was removed for fluoride measurement, and fresh water was placed in the sample vial. The aliquots were measured on an Orion Model 720A pH meter, using an Orion F⁻ detector, Model 96-09. Using a fluoride standard solution, the meter was referenced at 1, 0.1, and 0.01 ppm. The ionic strength of each sample, and the standards, were adjusted with TISAB II buffer just prior to measurement. This method is similar to that used by other researchers (de Araujo, Garcia-Goody et al. 1996; Verbeeck, De Maeyer et al. 1998). Measurements were made on days 1, 2, 3, 4, 5, 7 and 14.

EXAMPLE 7

Fluoride Release Test. Two sets of samples were prepared for testing. In the first group, an 80:20 PMMA-MAA random copolymer was used with the following

glass contents: 0.00, 6.25, 14.29, 25.00, 33.33, 40.00, 45.45, 50.00, 60.53, 70.00 and 80.00 weight%. The second set includes the following PMMA-MAA random copolymers: 75:25, 80:20, 95:5 and 100:0, containing 0 weight% and 50 weight% glass. The copolymers were obtained from Polysciences, Inc., Warrington, Pennsylvania, the PMMA from Sigma Chemical Co., St. Louis, Missouri, and the glass from Specialty Glass Inc., Oldsmar, Florida, with a composition of 25.10 wt% SiO₂, 37.45 wt% Al₂O₃, and 37.45 wt% CaF₂. The glass particles are irregular in shape, range from 1-20 microns, and have no surface treatment. The glass was incorporated into the copolymer as described hereinabove in Example 1.

Five glass ionomer samples 1 cm x 1 cm x 0.05 cm were made using a high temperature compression system for all fourteen variations of the material, including the different glass contents and the different copolymers containing 50 wt% glass. Each sample was placed in a vial containing 5 ml ddH₂O. These samples were maintained on a shaker at 37°C for the duration of the experiment. Each day the water was removed for fluoride measurement, and fresh water was placed in the sample vial. The aliquots were measured on an Orion Model 720A pH meter, using an Orion F⁻ detector, Model 96-09. Using a fluoride standard solution, the meter was referenced at 1, 0.1, and 0.01 ppm. The ionic strength of each sample, and the standards, were adjusted with TISAB II buffer just prior to measurement. Measurements were made on days 1, 2, 3, 4, 5 and 7.

Discussion of Experimental Results

Effects of glass content on mechanical properties. Figure 1 shows the compressive modulus data for the glass filled 80:20 polymer. The elastic modulus is comparable to that of the best commercial glass ionomer materials available (O'Brien 1997). A sigmoidal trend is observed, with the modulus remaining nearly the same until a critical glass percentage is reached, at

which the modulus increases rapidly. At high glass content, another critical glass percentage is attained where additional glass no longer improves the modulus. It is believed that, upon optimization of the compositions of the present invention, even better modulus data may be obtained, thereby surpassing that of the best available commercial glass ionomer materials.

Similar to the compressive modulus data, the Vickers hardness of the material shows a sigmoidal trend with increased glass content (Figure 2). These values compare favorably to those of commercially available glass ionomers and dentin tooth structure (O'Brien 1997). As with the modulus data, it is believed that, upon optimization of the compositions of the present invention, even better hardness data may be obtained, thereby surpassing that of commercially available glass ionomer materials.

In the PMMA-MAA glass ionomer, the carboxyl group on the methacrylic acid unit is able to bind with Ca^{+2} and Al^{+3} ions from the glass. Some of these ions are presumably leached out of the glass during the reaction creating crosslinking between polymer chains, and also forming links between the ionomer and the glass itself. This produces a polymeric matrix filled with glass particles surrounded by ionic aggregates. Without being bound to any theory, it is believed that, by developing these ionic interactions between the glass particles and the polymer matrix, the mechanical properties of the material are increased.

When interpreting the strength data, it has been found in the present invention that it may be beneficial to consider several interactions. Some of the mechanical properties of the material may be attributed to a mechanical interaction between glass particles. The glass and polymer packing density may also play an important role in the analysis of the data at high glass contents. In addition, it has been found that three

chemical interactions may come into play: the polymer-glass ionic bonding, the polymer-polymer ionic bonding, and the polymer-polymer hydrogen bonding.

If the interactions mentioned above are considered in the compressive modulus data interpretation, it can be seen that the glass-glass mechanical contact appears to dominate the modulus trend. As more glass particles are added to the material, it becomes less pliable and therefore more resistant to being compressed. Modulus is related to very small deformation, so at low glass contents, the particles are not forced to touch each other, showing little change in the modulus. Once a critical glass content is reached, the modulus increases rapidly. At about 50 wt% glass, the material assumes a continuous glass structure with the particles touching, and the resistance to being compressed levels off. At high glass content the effect of packing density comes into consideration. Since the glass possesses a range of sizes, the packing is improved over particles of all one size. See, for example, Cross, M., W.H. Douglas, et al., "The Relationship Between Filler Loading and Particle Size Distribution in Composite Resin Technology," Journal of Dental Research **62** (7): 850-852 (1983).

At a certain packing density, it appears that additional glass does not increase the number of glass-glass particle contacts, thus no longer improving the mechanical properties of the material. Other researchers have seen a linear increase of modulus as the glass content is increased in a glass-filled PMMA composite material. See, for example, Soderholm, K.-J., "Relationship between compressive yield strength and filler fractions of PMMA composites," Acta Odontol. Scand. **40**: 145-150 (1982). However their data do not cover as wide of a glass range.

Figure 3 exhibits the yield strength data for the 80:20 copolymer with glass filler. There is a linear

increase in yield strength to approximately 60 wt% glass, where it then levels off. Unlike the modulus data, the yield strength at low glass content increases linearly starting at 0 wt% glass. It is believed that this is due to the fact that yield strength is a function of larger strain than the modulus, and the glass particles are forced to touch as the material yields. Other researchers have seen a linear increase in yield strength for a glass filled PMMA, for both silane treated and untreated glass. See, for example, Soderholm (1982); and Krause, W.R., S.-H. Park, et al., "Mechanical properties of BIS-GMA resin short glass fiber composites," Journal of Biomedical Materials Research 23: 1195-1211 (1989). It is believed that the packing density discussed above is most likely responsible for the bend in the yield strength curve at high glass content.

The toughness data is shown in Figure 4. The addition of glass improved the toughness of 80:20 PMMA-MAA at glass concentrations below 70 wt% glass. The peak appears around about 15-20 wt% glass.

Effects of copolymer composition on mechanical properties. Examination of Figure 5 shows the elastic modulus data for PMMA-MAA copolymers with 0 wt% and 50 wt% glass. An increase in the methacrylic acid component of the copolymer increases the modulus of the material, as does the addition of glass filler. The Vickers hardness data show a similar trend (See Figure 6).

In Figure 7, the yield strength data for the PMMA-MAA glass ionomers are displayed. There is an increase in yield strength with an increase in MAA monomers. There is also an increase in yield strength with the addition of 50 wt% glass. There is a significant divergence in the two sets of data at higher ionomer content. The linear increase in yield strength with increase in MAA content (0 wt% glass) is presumably due to a greater number of hydrogen bonds between the chains. It is believed that the fact that the glass

ionomers with 50 wt% glass have higher yield strengths than those of the corresponding copolymers (0 wt% glass) can be attributed to the glass-glass mechanical interactions. It is further believed that the divergence between 0 wt% and 50 wt% glass samples in the increasing rate (slope) of yield strength with MAA content shows the synergism of the glass filler and ionic interactions--both the ionic links between polymer chains caused by the leached ions and the ionic links between the polymer chains and the glass fragments.

The toughness data are shown in Figure 8. For MAA concentrations of 25%, 20%, and 0%, the addition of 50 wt% glass appeared to improve the toughness of the copolymer. The copolymer sample containing 5% MAA with 0 wt% glass appears to have a higher toughness than a similar sample with 50 wt% glass--this appeared to be an aberrant test result, and the present inventors plan to repeat this experiment.

Fluoride Release. The results of the fluoride release testing are shown in Figures 9 and 10. There is an initial burst release that tapers off with time. This profile is typical for glass ionomer materials. This is important for the initial blocking of caries development. Overall, the plot shows that with an increase in glass content, there is an increase in fluoride release, allowing for fine-tuning of the amount of fluoride released into the oral environment.

Examination of how the fluoride release is affected by copolymer composition shows that there is an increase in the release rate with a greater MAA content. The MAA component affects the material in two ways. It possesses the chemical unit which leaches the ions out of the glass. Therefore, a higher MAA content would indicate that more fluoride is available for release. Secondly, an increase in the MAA component makes the polymer slightly more susceptible to water attack. This would also create an increase in the fluoride release.

It is believed that, over a longer time, the unique ionomer composite composition of the present invention will advantageously provide continuous fluoride release.

As discussed hereinabove, the properties of the novel and unique ionomer composite composition of the present invention may easily be adjusted by altering the ratio of hydrophilic monomer to hydrophobic monomer in the copolymer, and/or by altering the glass content.

Bonding between acid-containing hydrophilic monomers and glass particles, along with monomer-monomer bonds are formed by acid leaching of cations from the glass particles, forming a strong interface between the filler and matrix. An added advantage of this ion leaching is that fluoride ions may also be released from the glass, if desired.

While preferred embodiments, forms and arrangements of parts of the invention have been described in detail, it will be apparent to those skilled in the art that the disclosed embodiments may be modified. Therefore, the foregoing description is to be considered exemplary rather than limiting, and the true scope of the invention is that defined by the following claims.

What is claimed is:

1 1. An ionomer composite composition having
2 improved physical properties, the composite consisting
3 essentially of:

4 a glass material containing at least one of
5 divalent cations and multivalent cations; and
6 at least one copolymer, comprising:

7 at least one hydrophilic monomer
8 containing acid functional groups adapted to react with
9 the at least one of divalent cations and multivalent
10 cations to form ionic crosslinks among polymer chains,
11 the hydrophilic monomer present in an amount sufficient
12 to impart a desired degree of aqueous solubility to the
13 copolymer; and

14 at least one hydrophobic monomer present
15 in an amount sufficient to impart a desired degree of
16 structural stability to the composite composition when
17 exposed to an aqueous environment.

1 2. The ionomer composite composition as
2 defined in claim 1 wherein the glass material further
3 contains fluoride, and wherein the composite composition
4 provides substantially continuous fluoride release.

1 3. The ionomer composite composition as
2 defined in claim 1 wherein the glass material is selected
3 from the group consisting of SiO_2 , Al_2O_3 , AlF_3 , CaF_2 ,
4 NaF , Na_3AlF_6 , AlPO_4 , and mixtures thereof.

1 4. The ionomer composite composition as
2 defined in claim 1 wherein a weight ratio of glass
3 material to copolymer ranges between about 10:90 and
4 about 90:10.

1 5. The ionomer composite composition as
2 defined in claim 4 wherein the weight ratio of glass
3 material to copolymer ranges between about 40:60 and
4 about 85:15.

1 6. The ionomer composite composition as
2 defined in claim 1 wherein the at least one hydrophilic
3 monomer is a monomer containing unsaturated carboxylic
4 acid.

1 7. The ionomer composite composition as
2 defined in claim 6 wherein the unsaturated carboxylic
3 acid is selected from the group consisting of: acrylic
4 acid, methacrylic acid, 4-vinylbenzoic acid, crotonic
5 acid, oleic acid, elaidic acid, itaconic acid, maleic
6 acid, fumaric acid, acetylenedicarboxylic acid,
7 tricarballylic acid, sorbic acid, linoleic acid,
8 linolenic acid, eicosapentenoic acid, anhydrides of the
9 acids, derivatives of the unsaturated carboxylic acids,
10 and mixtures thereof.

1 8. The ionomer composite composition as
2 defined in claim 7 wherein the hydrophilic monomer is
3 selected from the group consisting of organic acids
4 having at least one of sulfonic acid and phosphonic acid
5 replacement of the carboxyl group of the unsaturated
6 carboxylic acids, derivatives of the organic acids, and
7 mixtures thereof.

1 9. The ionomer composite composition as
2 defined in claim 1 wherein the at least one hydrophobic
3 monomer is unsaturated.

1 10. The ionomer composite composition as
2 defined in claim 9 wherein the at least one hydrophobic
3 monomer is selected from the group consisting of
4 acrylates, methacrylates, ethylenes, propylenes, tetra-
5 fluoroethylene, styrenes, vinyl chloride, vinylidene
6 chloride, vinyl acetate, acrylonitrile, 2,2-bis[4-(2-
7 hydroxy-3-methacryloyloxy-propoxy)-phenyl] propane
8 (Bis-GMA), ethyleneglycol dimethacrylate (EGDMA), tri-
9 ethyleneglycol dimethacrylate (TEGDMA), bis(2-methacryly-
10 oxyethyl) ester of isophthalic acid (MEI), bis(2-meth-

11 acrylyoxyethyl) ester of terephthalic acid (MET), bis(2-
12 methacrylyoxyethyl) ester of phthalic acid (MEP), 2,2-
13 bis(4-methacrylyoxy phenyl) propane (BisMA), 2,2-bis[4-
14 (2-methacrylyloxyethoxy) phenyl] propane (BisEMA), 2,2-
15 bis[4-(3-methacrylyloxy-propoxy) phenyl] propane
16 (BisPMA), hexafluoro-1,5-pentanediol dimethacrylate
17 (HFPDMA), bis-(2-methacrylyloxyethoxy-hexafluoro-2-
18 propyl) benzene [Bis(MEHFP) ϕ], 1,6-bis(methacrylyloxy-2-
19 ethoxycarbonylamino)-2,4,4-tri-methylhexan (UEDMA), spiro
20 orthocarbonates, derivatives of the hydrophobic monomers,
21 and mixtures thereof.

1 11. The ionomer composite composition as
2 defined in claim 1 wherein a ratio of the hydrophilic
3 monomer to the hydrophobic monomer is between about 1:99
4 and about 99:1.

1 12. The ionomer composite composition as
2 defined in claim 1 wherein the at least one copolymer is
3 selected from the group consisting of poly(methyl
4 methacrylate-methacrylic acid), poly(methyl acrylate-
5 acrylic acid), poly(methyl methacrylate-acrylic acid),
6 poly(ethyl acrylate-acrylic acid), poly(ethyl
7 methacrylate-methacrylic acid), poly(butyl acrylate-
8 acrylic acid), poly(ethylene-acrylic acid),
9 poly(ethylene-methacrylic acid), poly(acrylonitrile-
10 maleic anhydride), poly(butadiene-acrylonitrile-acrylic
11 acid), poly(butadiene-maleic acid), poly(butadiene-maleic
12 anhydride), poly(acrylamide-acrylic acid), poly(2-
13 hydroxyethyl methacrylate-methacrylic acid),
14 poly(propylene-acrylic acid), poly(propylene-ethylene-
15 acrylic acid), poly(vinyl chloride-vinyl acetate-maleic
16 acid), and mixtures thereof.

1 13. The ionomer composite composition as
2 defined in claim 1 wherein the composite further includes
3 as a minor constituent a polymer selected from the group

4 consisting of PMMA, polycarbonates, polyethylenes,
5 polyamides, PEEK polymers, epoxies, and mixtures thereof.

1 14. The ionomer composite composition as
2 defined in claim 2 wherein the composite further includes
3 a minor constituent selected from the group consisting of
4 pigments, surfactants, adhesion enhancers, fluoride
5 releasing enhancers, bioactive agents, and mixtures
6 thereof.

1 15. The ionomer composite composition as
2 defined in claim 2 wherein the glass material consists of
3 25.10 wt% SiO₂, 37.45 wt% Al₂O₃, and 37.45 wt% CaF₂.

1 16. The ionomer composite composition as
2 defined in claim 1 wherein the at least one copolymer is
3 poly(methyl methacrylate-co-methacrylic acid) (PMMA-MAA).

1 17. The ionomer composite composition as
2 defined in claim 16 wherein the ratio of PMMA to MAA is
3 80:20.

1 18. An ionomer composite composition having
2 improved physical properties and useful for dental
3 applications, the composite consisting essentially of:

4 a glass material containing at least one of
5 divalent cations and multivalent cations; and

6 at least one copolymer, comprising:

7 at least one hydrophilic monomer

8 containing acid functional groups adapted to react with
9 the at least one of divalent cations and multivalent
10 cations to form ionic crosslinks among polymer chains,
11 the hydrophilic monomer present in an amount sufficient
12 to impart a desired degree of aqueous solubility to the
13 copolymer; and

14 at least one hydrophobic monomer present
15 in an amount sufficient to impart a desired degree of

16 structural stability to the composite composition when
17 exposed to an aqueous environment;

18 wherein the composite composition is adaptable
19 for use in dental applications ranging from those
20 requiring water soluble compositions, to intermediary
21 materials, to those requiring water insoluble
22 compositions.

1 19. The ionomer composite composition as
2 defined in claim 18 wherein the glass material further
3 contains fluoride, and wherein the composite composition
4 provides substantially continuous fluoride release.

1 20. The ionomer composite composition as
2 defined in claim 19 wherein the glass material is
3 selected from the group consisting of SiO_2 , Al_2O_3 , AlF_3 ,
4 CaF_2 , NaF , Na_3AlF_6 , AlPO_4 , and mixtures thereof.

1 21. The ionomer composite composition as
2 defined in claim 20 wherein the weight ratio of glass
3 material to copolymer ranges between about 40:60 and
4 about 85:15.

1 22. The ionomer composite composition as
2 defined in claim 21 wherein the at least one hydrophilic
3 monomer is an unsaturated carboxylic acid selected from
4 the group consisting of: acrylic acid, methacrylic acid,
5 4-vinylbenzoic acid, crotonic acid, oleic acid, elaidic
6 acid, itaconic acid, maleic acid, fumaric acid,
7 acetylenedicarboxylic acid, tricarballylic acid, sorbic
8 acid, linoleic acid, linolenic acid, eicosapentenoic
9 acid, anhydrides of the acids, derivatives of the
10 unsaturated carboxylic acids, and mixtures thereof.

1 23. The ionomer composite composition as
2 defined in claim 22 wherein the at least one hydrophilic
3 monomer is selected from the group consisting of organic
4 acids having at least one of sulfonic acid and phosphonic

5 acid replacement of the carboxyl group of the unsaturated
6 carboxylic acids, derivatives of the organic acids, and
7 mixtures thereof.

1 24. The ionomer composite composition as
2 defined in claim 22 wherein the at least one hydrophobic
3 monomer is unsaturated and is selected from the group
4 consisting of acrylates, methacrylates, ethylenes,
5 propylenes, tetra-fluoroethylene, styrenes, vinyl
6 chloride, vinylidene chloride, vinyl acetate,
7 acrylonitrile, 2,2-bis[4-(2-hydroxy-3-methacryloyloxy-
8 propyloxy)-phenyl] propane (Bis-GMA), ethyleneglycol
9 dimethacrylate (EGDMA), tri-ethyleneglycol dimethacrylate
10 (TEGDMA), bis(2-methacryly-oxyethyl) ester of isophthalic
11 acid (MEI), bis(2-meth-acrylyoxyethyl) ester of
12 terephthalic acid (MET), bis(2-methacrylyoxyethyl) ester
13 of phthalic acid (MEP), 2,2-bis(4-methacrylyoxy phenyl)
14 propane (BisMA), 2,2-bis[4-(2-methacrylyloxyethoxy)
15 phenyl] propane (BisEMA), 2,2-bis[4-(3-methacrylyloxy-
16 propoxy) phenyl] propane (BisPMA), hexafluoro-1,5-
17 pentanediol dimethacrylate (HFPDMA), bis-(2-
18 methacrylyloxyethoxy-hexafluoro-2-propyl) benzene
19 [Bis(MEHFP) ϕ], 1,6-bis(methacrylyloxy-2-
20 ethoxycarbonylamino)-2,4,4-tri-methylhexan (UEDMA), spiro
21 orthocarbonates, derivatives of the hydrophobic monomers,
22 and mixtures thereof.

1 25. The ionomer composite composition as
2 defined in claim 24 wherein a ratio of the hydrophilic
3 monomer to the hydrophobic monomer is between about 1:99
4 and about 99:1.

1 26. The ionomer composite composition as
2 defined in claim 18 wherein the at least one copolymer is
3 selected from the group consisting of poly(methyl
4 methacrylate-methacrylic acid), poly(methyl acrylate-
5 acrylic acid), poly(methyl methacrylate-acrylic acid),
6 poly(ethyl acrylate-acrylic acid), poly(ethyl

7 methacrylate-methacrylic acid), poly(butyl acrylate-
8 acrylic acid), poly(ethylene-acrylic acid),
9 poly(ethylene-methacrylic acid), poly(acrylonitrile-
10 maleic anhydride), poly(butadiene-acrylonitrile-acrylic
11 acid), poly(butadiene-maleic acid), poly(butadiene-maleic
12 anhydride), poly(acrylamide-acrylic acid), poly(2-
13 hydroxyethyl methacrylate-methacrylic acid),
14 poly(propylene-acrylic acid), poly(propylene-ethylene-
15 acrylic acid), poly(vinyl chloride-vinyl acetate-maleic
16 acid), and mixtures thereof.

1 27. The ionomer composite composition as
2 defined in claim 18 wherein the composite further
3 includes as a minor constituent a polymer selected from
4 the group consisting of PMMA, polycarbonates,
5 polyethylenes, polyamides, PEEK polymers, epoxies, and
6 mixtures thereof.

1 28. The ionomer composite composition as
2 defined in claim 19 wherein the composite further
3 includes a minor constituent selected from the group
4 consisting of pigments, surfactants, adhesion enhancers,
5 fluoride releasing enhancers, bioactive agents, and
6 mixtures thereof.

1 29. The ionomer composite composition as
2 defined in claim 19 wherein the glass material consists
3 of 25.10 wt% SiO₂, 37.45 wt% Al₂O₃, and 37.45 wt% CaF₂.

1 30. The ionomer composite composition as
2 defined in claim 19 wherein the at least one copolymer is
3 poly(methyl methacrylate-co-methacrylic acid) (PMMA-MAA).

1 31. The ionomer composite composition as
2 defined in claim 30 wherein the ratio of PMMA to MAA is
3 80:20.

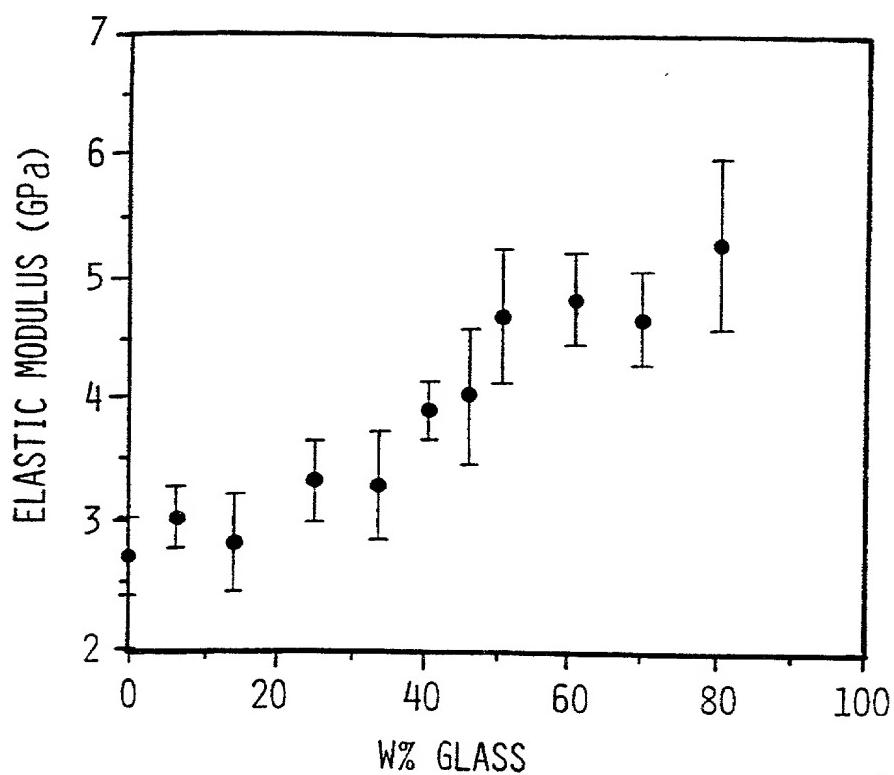


FIG. 1

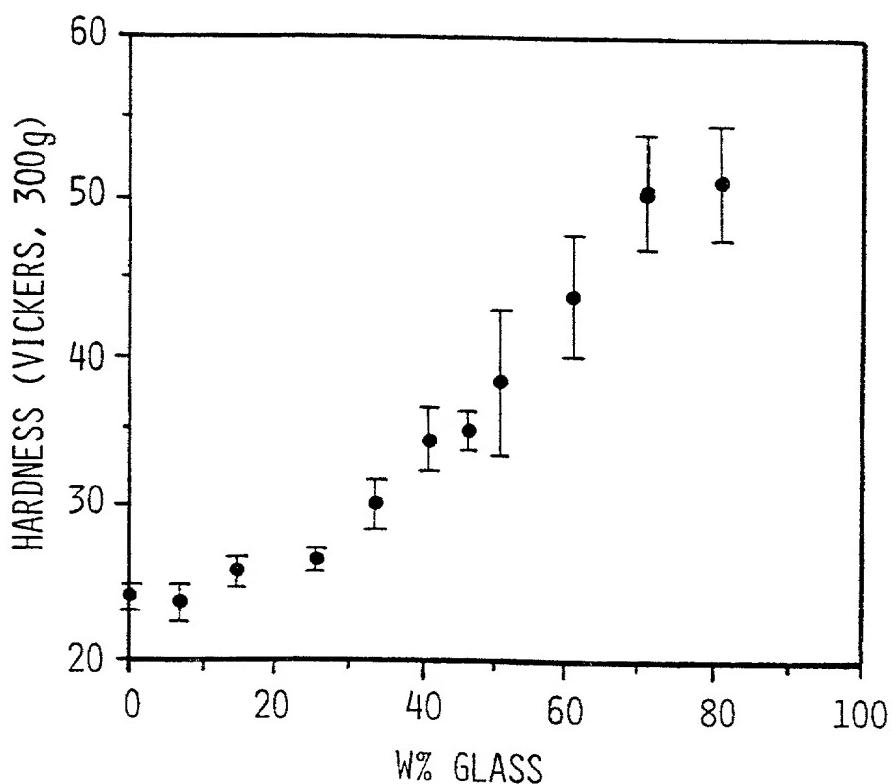


FIG. 2

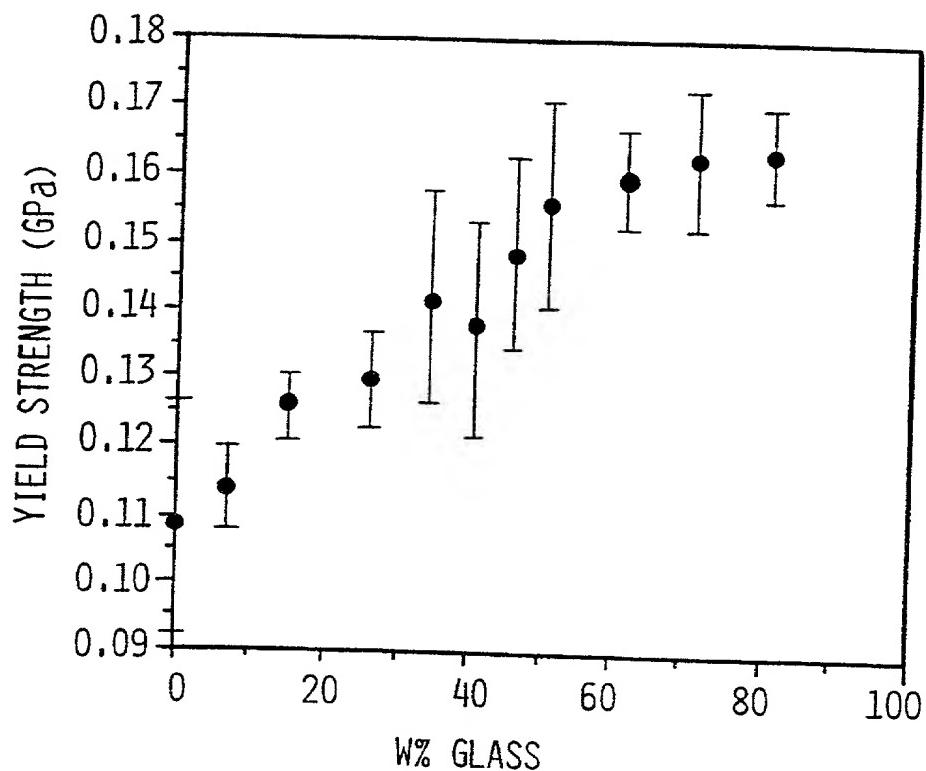


FIG. 3

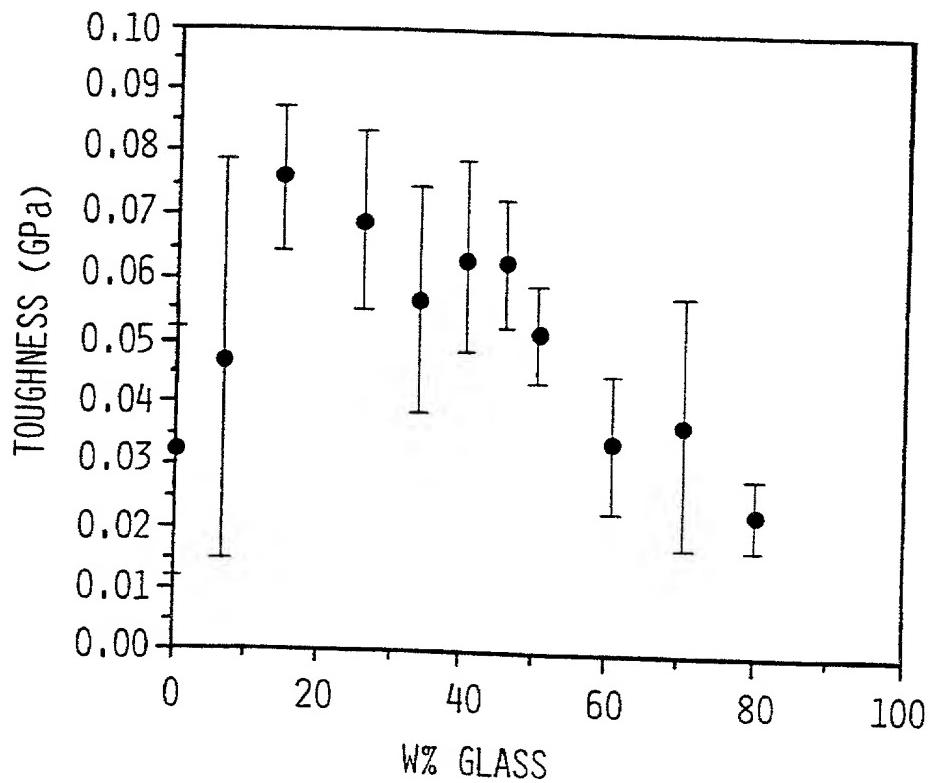


FIG. 4

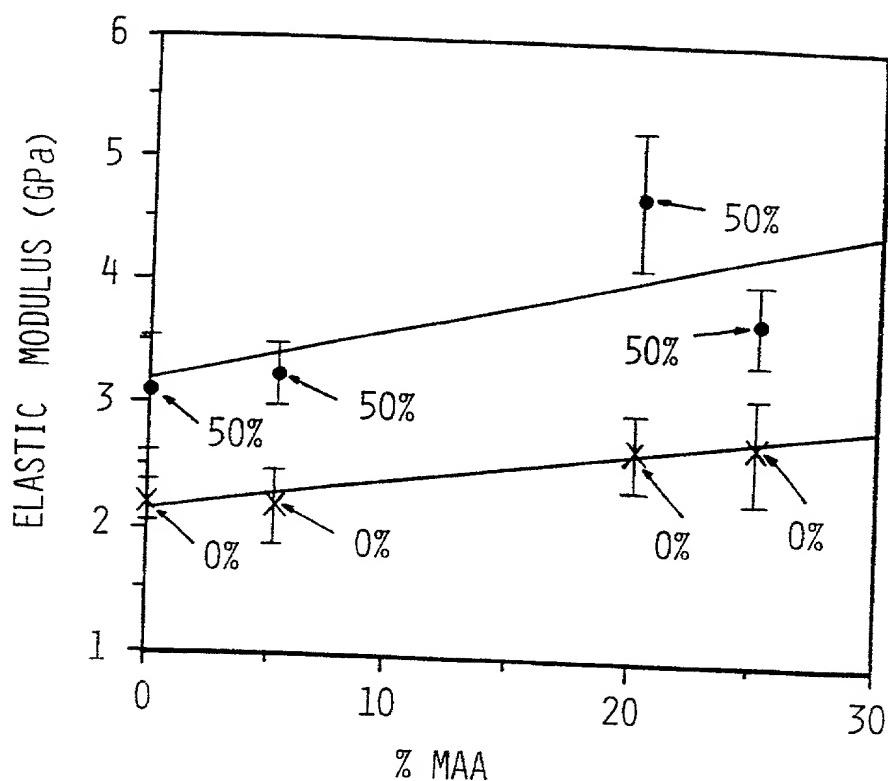


FIG. 5

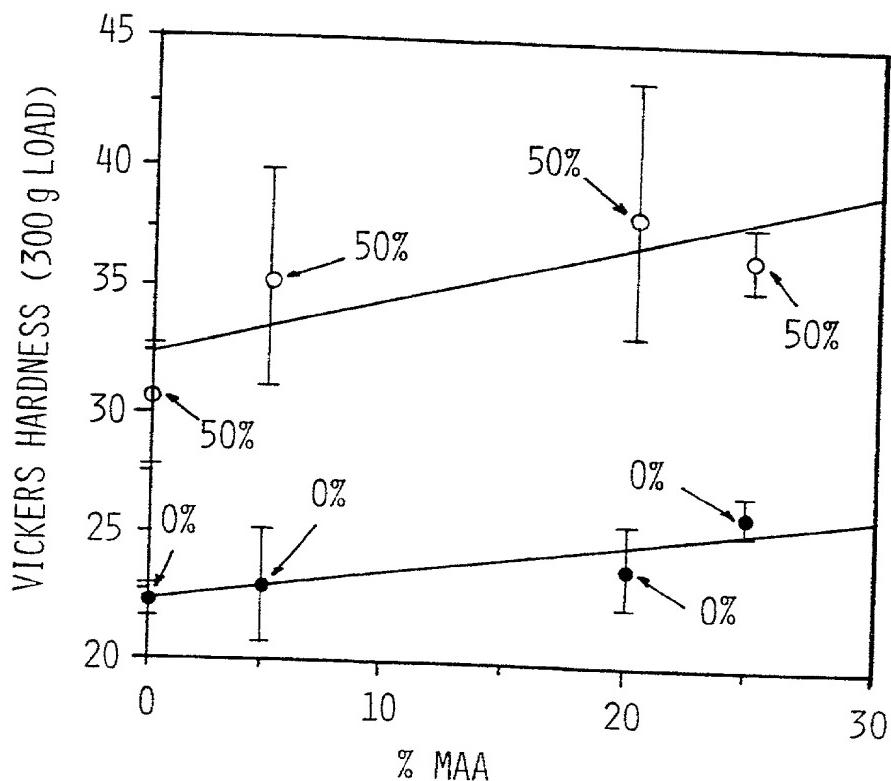


FIG. 6

4 / 6

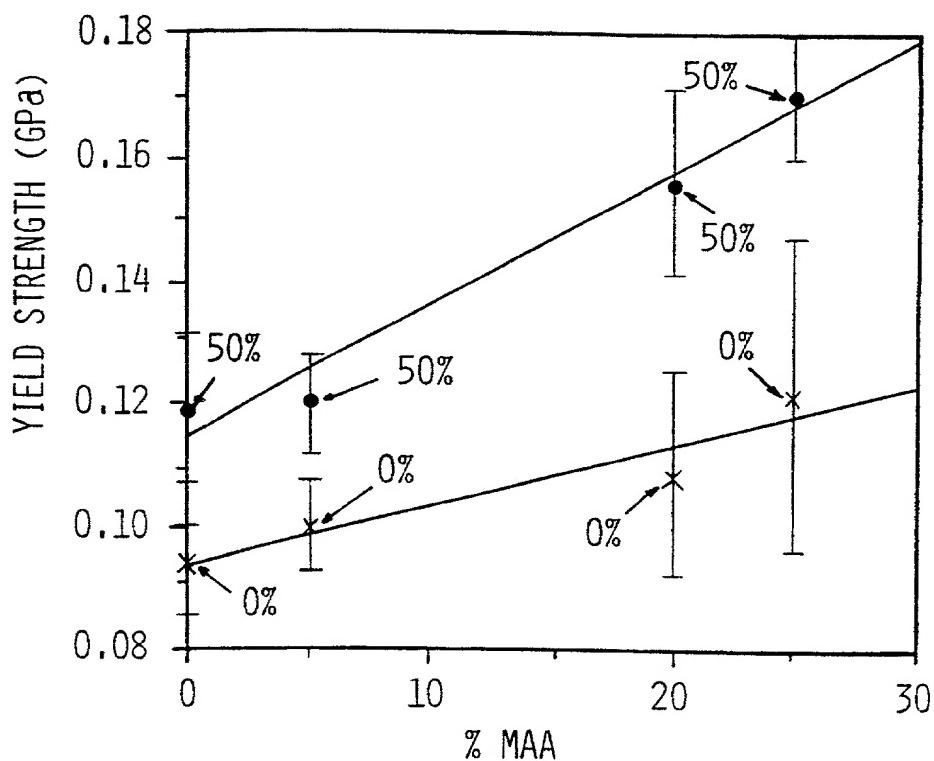


FIG 7

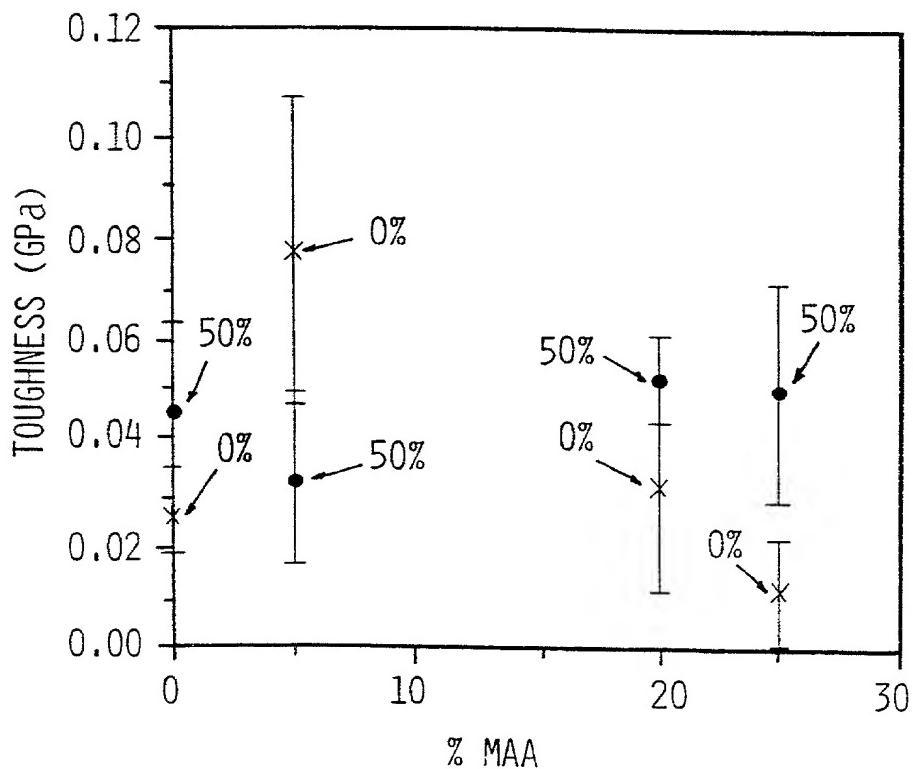


FIG 8

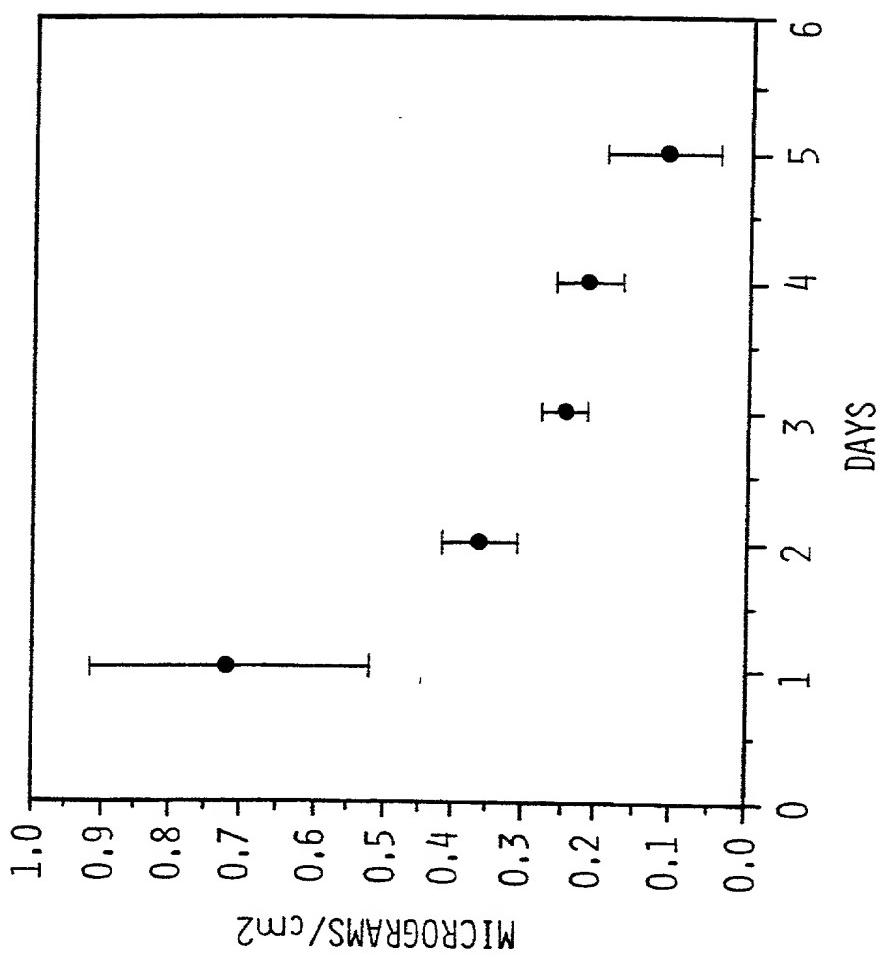


FIG. 9

09/936692

WO 00/55253

PCT/US00/06974

6 / 6

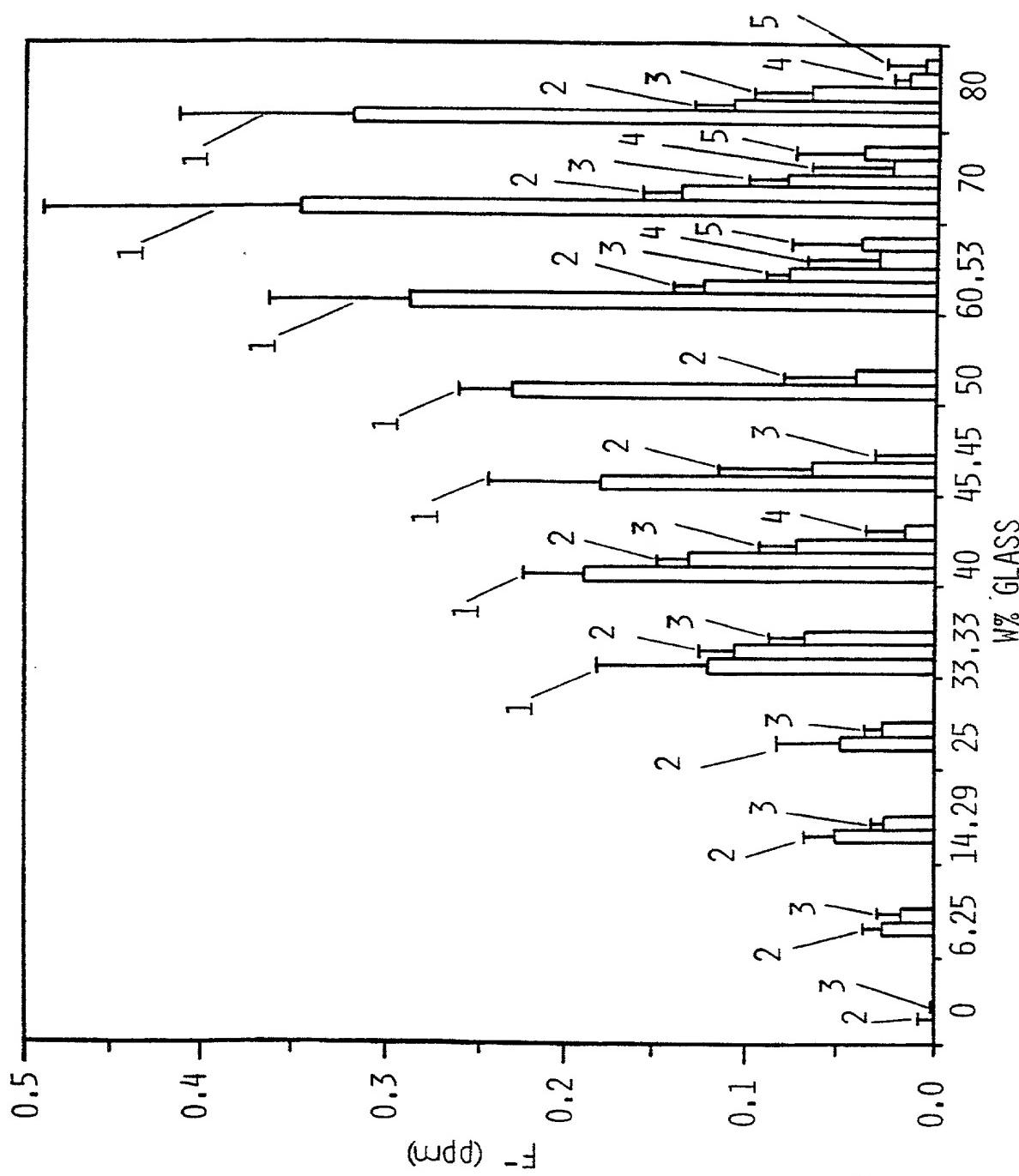


FIG. 10

POWER OF ATTORNEY:

I hereby appoint the following attorney(s) and/or agent(s) Julia Church Dierker, Patent Office Registration No. 33368, Andrew R. Basile, Patent Office Registration No. 24753, William M. Hanlon, Jr., Patent Office Registration No. 28422, and Thomas D. Helmholdt, Patent Office Registration No. 33181 as my attorney(s) and/or agent(s), to prosecute this application and to transact all business in the United States Patent and Trademark Office connected therewith.

Send all correspondence to: **Julia Church Dierker**
Young & Basile, P.C.
3001 West Big Beaver Road, Suite 624
Troy, Michigan 48084
Phone: (248) 649-3333

I hereby declare that all statements made herein of my own knowledge are true and that all statements made on information and belief are believed to be true; and further that these statements were made with the knowledge that willful false statements and the like so made are punishable by fine or imprisonment, or both, under §1001 of Title 18 of the United States Code and that such willful false statements may jeopardize the validity of the application or any patent issued thereon.

Full name of sole or first inventor Peter X. Ma

Inventor's Signature SEE SEPARATE DOCUMENT

Date _____ Citizenship USA

Residence 3208 Foxway Court, Ann Arbor, Michigan 48105

Post Office Address (Same as above)

Full name of second joint inventor, if any Anne E. Huber

Inventor's Signature Anne E. Huber

Date 8-21-01 Citizenship USA

Residence 44115 Stassen, Novi, Michigan 48375

Post Office Address (Same as above)

Full name of third joint inventor, if any _____

Inventor's Signature _____

Date _____ Citizenship _____

Residence _____

Post Office Address (Same as above)

Full name of fourth joint inventor, if any _____

Inventor's Signature _____

Date _____ Citizenship _____

Residence _____

Post Office Address (Same as above)

Our Reference: UMJ-105-B (UM-1667)

COMBINED DECLARATION AND POWER OF ATTORNEY**DECLARATION:**

As a below named inventor, I hereby declare that:

My residence, post office address and citizenship are as stated below next to my name.

I believe I am the original, first and sole inventor (if only one name is listed below) or an original, first and joint inventor (if plural names are listed below) of the subject matter which is claimed and for which a patent is sought on the invention entitled:

IONOMER COMPOSITE COMPOSITIONS

the specification of which (check only one item below):

[] is attached hereto.

[] was filed as United States application Serial No. _____ on _____, and was amended on or through _____ (if applicable).

[X] was filed as PCT international application Number PCT/US00/06974 on March 16, 2000, and was amended under PCT Article 19 on _____ (if applicable).

I hereby state that I have reviewed and understand the contents of the above identified specification, including the claims, as amended by any amendment referred to above.

I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, §1.56.

I hereby claim foreign priority benefits under Title 35, United States Code, §119(a)-(d) or §365(b) of any foreign application(s) for patent or inventor's certificate or §365(a) of any PCT international application(s) which designated at least one country other than the United States of America, listed below and have also identified below, by checking the box, any foreign application for patent or inventor's certificate, or PCT international application(s) having a filing date before that of the application on which priority is claimed:

Prior Foreign/PCT Application(s) and any Priority Claims Under 35 U.S.C. §119:			Priority Claimed	
(Number)	(Country)	(Day/Mo/Yr Filed)	[X]	[]
(Number)	(Country)	(Day/Mo/Yr Filed)	Yes	No
			[]	[]
			Yes	No

I hereby claim the benefit under 35 U.S.C. §119(e) of any United States provisional application(s) listed below.

(Application Number)	(Filing Date)
60/124,819	March 17, 1999

I hereby claim the benefit under Title 35, United States Code, §120 of any United States application(s) or §365(c) of any PCT international application(s) designating the United States of America, listed below and, insofar as the subject matter of each of the claims of this application is not disclosed in the prior United States or PCT international application(s) in the manner provided by the first paragraph of Title 35, United States Code, §112, I acknowledge the duty to disclose information which is material to patentability as defined in Title 37, Code of Federal Regulations, §1.56 which became available between the filing date of the prior application and the national or PCT international filing date of this application.

Prior U. S. Application(s) or PCT International Application(s) Designating the U.S. for Benefit Under 35 U.S.C. §120:

(Application Number)	(Filing Date)	(Status: patented, pending, abandoned)
(Application Number)	(Filing Date)	(Status: patented, pending, abandoned)